Opportunity for Human Research Subjects

Treatment Study for Reducing Symptoms of Post-Traumatic Stress

Category Enrollment Closing Date
PTSD/TBI Oct 1, 2018

Description
This is a 12-week study investigating the use of TNX-102 SL—a sublingual formulation of cyclobenzaprine*. Study participation includes 5 in-clinic visits and 1 follow-up phone call over approximately 13-17 weeks. Compensation up to $375 is available for completion of study.

*the same active ingredient as Flexeril®

Requirements
You may be eligible if you are a Veteran who has experienced stress, anxiety, or insomnia due to a military-related traumatic event after 2001.

Benefits
Your participation may help to inform treatment options for veterans with post-traumatic stress.

Contact Information

Name  Study Coordinator
Phone  (858) 552 - 8585  Extension: 7929

VA San Diego Healthcare System
IRB NUMBER: H160178
IRB APPROVAL DATE: 03/12/2018